



Patent Docket P1089R1C1

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of Lam et al. Serial No.: 09/724,868 Filed: 28 November 2000 For: ANTIBODY FORMULATION	Group Art Unit: Unassigned Examiner: Unassigned <div style="border: 1px solid black; padding: 5px;"><p style="text-align: center;">CERTIFICATE OF MAILING</p><p>I hereby certify that this correspondence is being deposited with the United States Postal Service with sufficient postage as first class mail in an envelope addressed to: Assistant Commissioner of Patents, Washington, D.C. 20231 on</p><p style="text-align: right;">February 13, 2001</p><p style="text-align: right;"><i>Ann Savelli</i></p><p style="text-align: right;">Ann Savelli</p></div>
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INFORMATION DISCLOSURE STATEMENT

Assistant Commissioner of Patents
Washington, D.C. 20231

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Sir:

Applicants submit herewith patents, publications or other information (attached hereto and listed on the attached revised Form PTO-1449) of which they are aware, which they believe may be material to the examination of this application and in respect of which there may be a duty to disclose in accordance with 37 CFR §1.56.

This Information Disclosure Statement is filed in accordance with the provisions of:

☒ 37 CFR §1.97(b)

- within three months of the filing date of the application other than a continued prosecution application under 37 CFR §1.53(d); or
- within three months of the date of entry of the national stage of a PCT application as set forth in 37 CFR §1.491, or
- before the mailing of the first Office action on the merits; or
- before the mailing of the first Office action after the filing of a request for a continued examination under 37 CFR §1.114.

☐ 37 CFR §1.97(c)

- by the applicant after the period specified in 37 CFR §1.97(b), but prior to the mailing date of any of a final action under 37 CFR §1.113, or a notice of allowance under 37 CFR §1.311, or an action that otherwise closes prosecution in the application, and is accompanied by either the fee set forth in 37 CFR §1.17(p) or a statement as specified in 37 CFR §1.97(e), as checked below.

☐ 37 CFR §1.97(d)

- after the period specified in CFR §1.97(c), and is accompanied by the fee set forth in 37 CFR §1.17(p) and a statement as specified in 37 CFR §1.97(e), as checked below.

02/21/2001 TGEDAMU1 00000071 070630 09724868

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[If either of boxes 37 CFR §1.97(c) or 37 CFR §1.97(d) is checked above, the following statement under 37 CFR §1.97(e) may need to be completed.]

- ☐ **37 CFR §1.97(e)** Each item of information contained in the information disclosure statement was first cited in any communication from a foreign patent office in a counterpart foreign application not more than three months prior to the filing of this information disclosure statement.
- ☐ **37 CFR §1.704(d)** Each item of information contained in the information disclosure statement was cited in a communication from a foreign patent office in a counterpart foreign application and the communication was not received by any individual designated in §1.56(c) more than thirty days prior to the filing of this information disclosure statement. Therefore, in accordance with the provisions of 37 CFR §1.704(d), the filing of this information disclosure statement will not be considered a failure to engage in reasonable efforts to conclude prosecution under 37 CFR §1.704.
- ☐ The U.S. Patent and Trademark Office is hereby authorized to charge Deposit Account No. 07-0630 in the amount of \$180.00 to cover the cost of this Information Disclosure Statement under 37 CFR §1.17(p). Any deficiency or overpayment should be charged or credited to this deposit account.

A list of the patent(s) or publication(s) is set forth on the attached revised Form PTO-1449 (Modified).

Those patent(s) or publication(s) which are marked with an asterisk (*) in the attached PTO-1449 form are not supplied because they were previously cited by or submitted to the Office in a prior application Serial No. 09/097,171, filed June 6, 1998 and relied upon in this application for an earlier filing date under 35 USC §120.

☐ BLAST results enclosed:

The undersigned also wishes to bring to the attention of the Examiner BLAST results of computerized alignments of the against sequences contained in the nucleotide and protein databases. The BLAST results are provided in paper form and are identified as reference "BLAST Results A-1- A-()" (nucleotide) and "BLAST Results B-1 - B-()" (protein) on the PTO Form 1449. Applicant requests that these references also be considered and that the Form 1449 be initialed to indicate the Examiner's consideration of the references.

A concise explanation of relevance of the items listed on PTO-1449 is:

- ☒ not given
- ☐ given for each listed item
- ☐ given for only non-English language listed item(s) [Required]

- ☐ in the form of an English language copy of a Search Report from a foreign patent office, issued in a counterpart application, which refers to the relevant portions of the references.

In accordance with 37 CFR §1.97(g), the filing of this information disclosure statement shall not be construed as a representation that a search has been made.

In accordance with 37 CFR §1.97(h), the filing of this information disclosure statement shall not be construed to be an admission that the information cited in the statement is, or is considered to be, material to patentability as defined in 37 CFR § 1.56(b).

In the event that the Office determines a fee to be due where none is specifically authorized in this paper, the U.S. Patent and Trademark Office is hereby authorized to charge Deposit Account No. 07-0630 in the amount of \$180.00 to cover the cost of this Information Disclosure Statement under 37 CFR §1.17(p).

Respectfully submitted,

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Telephone No. (650) 225-4462

Date: February 13, 2001



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FORM PTO-1449

U.S. Dept. of Commerce
Patent and Trademark OfficeAtty Docket No.
P1089R1C1Serial No.
09/724,868

LIST OF DISCLOSURES CITED BY APPLICANT

(Use several sheets if necessary)

Applicant
Lam et al.Filing Date
28 Nov 2000Group
Unassigned

U.S. PATENT DOCUMENTS

Examiner Initials		Document Number	Date	Name	Class	Subclass	Filing Date
	* 1	4,093,606	06.06.78	Coval, M.			
	* 2	4,457,916	03.07.84	Hayashi et al.			
	* 3	4,499,073	12.02.85	Tenold, R.			
	* 4	4,877,608	31.10.89	Lee, T. et al.			
	* 5	4,940,782	10.07.90	Rup et al.			
	* 6	5,032,405	16.07.91	Huang, H. et al.			
	* 7	5,036,049	30.07.91	Audhya, T. et al.			
	* 8	5,096,885	17.03.92	Pearlman et al.			
	* 9	5,147,637	15.09.92	Wright et al.			
	*10	5,149,653	22.09.92	Roser			
	*11	5,215,743	01.06.93	Singh, M. et al.			
	*12	5,262,296	16.11.93	Ogawa, E. et al.			
	*13	5,307,640	03.05.94	Fawzy et al.			
	*14	5,399,670	21.03.95	Bhattacharya, P. et al.			
	*15	5,506,342	09.04.96	Reno et al.			
	*16	5,580,856	03.12.96	Prestrelski et al.			
	*17	5,589,167	31.12.96	Cleland et al.			
	*18	5,608,038	04.03.97	Eibl et al.			
	*19	5,654,403	05.08.97	Smith et al.			
	*20	5,730,980	24.03.98	Ulevitch et al.			
	*21	5,736,137	07.04.98	Anderson et al.			
	*22	5,770,700	23.06.98	Webb et al.			

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FOREIGN PATENT DOCUMENTS

Examiner Initials		Document Number	Date	Country	Class	Subclass	Translation Yes No	
	*23	AU-B-30771/89	07.09.89	AUSTRALIA				
	*24	2,138,853	29.06.95	CANADA				
	*25	303,746	22.02.89	EPO				
	*26	391,444	10.10.90	EPO				
	*27	661,060	05.07.95	EPO (WITH ENGLISH ABSTRACT)				
	*28	WO 89/09402	05.10.89	PCT (WITH ENGLISH ABSTRACT)				
	*29	WO 89/11297	30.11.89	PCT				
	*30	WO 90/11091	04.10.90	PCT				
	*31	WO 92/22653	23.12.92	PCT				
	*32	WO 94/11026	26.05.94	PCT				
	*33	WO 94/26302	24.11.94	PCT				

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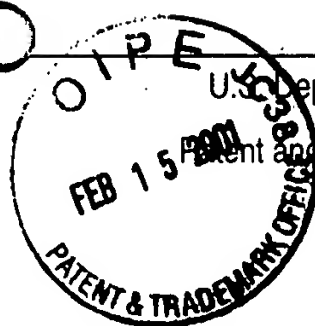
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	*34	WO 96/41164	19.12.96	PCT				
	*35	WO 97/04801	13.02.97	PCT				
	*36	WO 97/04807	13.02.97	PCT				
	*37	WO 97/17087	15.05.97	PCT				

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	*38	Akers, M., "Considerations in selecting antimicrobial preservative agents for parenteral product development" <u>Pharmaceutical Technology</u> pps. 36-40;43-44;46 (May 1984)						
	*39	Albelda et al., "Integrins and other cell adhesion molecules" <u>FASEB-J</u> 4(11):2868-2880 (1990)						
	*40	Arakawa et al., "Protein-Solvent Interactions in Pharmaceutical Formulations" <u>Pharmaceutical Research</u> 8(3):285-291 (1991)						
	*44	Bam et al., "Stability of Protein Formulations: Investigation of Surfactant Effects by a Novel EPR Spectroscopic Technique" <u>Pharm. Res.</u> 12:2-11 (1995)						
	*42	Beauvais et al., "Both Glassy State and Native Structure are Required for Storage Stability of Lyophilized Interleukin-1 Receptor Antagonist" <u>Pharm. Res.</u> (Abstract #2007) 12(9):S-80 (1995)						
	*40	Bogard et al., "Practical considerations in the production, purification, and formulation of monoclonal antibodies for immunoscintigraphy and immunotherapy" <u>Seminars in Nuclear Medicine</u> 19(3):202-220 (1989)						
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	*49	Chang et al., "Development of a Stable Freeze-dried Formulation of Recombinant Human Interleukin-1 Receptor Antagonist" <u>Pharmaceutical Research</u> 13(2):243-248 (1996)						
	*46	Chang et al., "Nucleotide sequence of the alkaline phosphatase gene of Escherichia Coli" <u>Gene</u> 44:121-125 (1986)						
	*47	Clarke et al., "Lability of Asparagine and Aspartic Acid Residues in Proteins and Peptides" <u>Stability of Protein Pharmaceuticals. Part A: Chemical and Physical Pathways of Protein Degradation</u> , T.J. Ahern and M.C. Manning, New York:Plenum Press, Chapter 1, pps. 1-29. (1992)						
	*48	Cleland and Jones, "Development of Stable Protein Formulations for Microencapsulation in Biodegradable Polymers" <u>Proceed. Intern. Symp. Control. Rel. Bioact. Mater.</u> 22:514-515 (1995)						
	*49	Cleland et al., "Mechanisms of Nonionic Surfactant Stabilization of Proteins" <u>Pharmaceutical Research</u> (Abstract #BIOTEC 2012; Ninth Annual Meeting of the American Association of Pharmaceutical Scientists held in San Diego, CA on November 6-10, 1994) 11(10 Suppl.):S73 (1994)						
	*50	Cleland et al., "The Development of Stable Protein Formulations: A Close Look at Protein Aggregation, Deamidation, and Oxidation" <u>Critical Reviews in Therapeutic Drug Carrier Systems</u> 10(4):307-377 (1993)						
	*51	Draber et al., "Stability of Monoclonal IgM Antibodies Freeze-Dried in the Presence of Trehalose" <u>Journal of Immunological Methods</u> 181(1):37-43 (1995)						
	*52	Hernandez et al., "Role of neutrophils in ischemia-reperfusion-induced microvascular injury" <u>Am. J. Physiol.</u> 253(3 Pt 2):H699-H703 (1987)						
	*53	Hildreth et al., "A Human Lymphocyte-associated Antigen Involved in Cell-mediated Lympholysis" <u>European Journal of Immunology</u> 13:202-208 (1983)						

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*54	Hildreth et al., "The human lymphocyte function-associated (HLFA) antigen and a related macrophage differentiation antigen (HMac-1): functional effects of subunit-specific monoclonal antibodies" <u>J. Immunol.</u> 134:3272-3280 (1985)
*55	Hynes, "Integrins: versatility, modulation, and signaling in cell adhesion" <u>Cell</u> 69(1):11-25 (1992)
*56	Izutsu et al., "The effects of additives on the stability of freeze-dried β -galactosidase stored at elevated temperature" <u>Intl. J. Pharmaceutics</u> 71:137-146 (1991)
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*58	Kossiakoff, A.A., "Tertiary Structure Is a Principal Determinant to Protein Deamidation" <u>Science</u> 240:191-194 (1988)
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*60	Lam et al., "Microencapsulation of Recombinant Humanized Monoclonal Antibody for Local Delivery" <u>Proceedings of the 24th International Symposium on Controlled Release of Bioactive Materials</u> 24:759-760 (June 15-19, 1997)
*61	Lam et al., "Pitfalls in Development of Multi-Dose Liquid Formulations for Three Protein Pharmaceuticals" <u>ACS National Meeting, New Orleans</u> (Abstract #137 and slides presented) (March 24-28, 1996)
*62	Li et al., "Aggregation and Precipitation of Human Relaxin Induced by Metal-Catalyzed Oxidation" <u>Biochemistry</u> 34(17):5762-5772 (1995)
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*64	Mileski et al., "Inhibition of CD18-dependent neutrophil adherence reduces organ injury after hemorrhagic shock in primates" <u>Surgery</u> 108:206-212 (1990)
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*67	Pearlman et al., "Analysis of Protein Drugs" <u>Peptide and Protein Drug Delivery</u> , Vincent H. L. Lee, Marcel Dekker, Inc., Chapter 6, pps. 247-301 (1991)
*68	Picken et al., "Nucleotide sequence of the gene for heat-stable enterotoxin II of Escherichia coli" <u>Infection and Immunity</u> 42(1):269-275 (1983)
*69	Pikal et al., "The Effects of Formulation Variables on the Stability of Freeze-Dried Human Growth Hormone" <u>Pharm. Res.</u> 8:427-436 (1991)
*70	Pikal, M., "Freeze-Drying of Proteins, Part 2: Formulation Selection" <u>Biopharm.</u> 3(9):26-30 (1990)
*71	Rao and Kroon, "Orthoclone OKT3: Chemical Mechanisms and Functional Effects of Degradation of a Therapeutic Monoclonal Antibody" <u>Stability and Characterization of Protein and Peptide Drugs: Case Histories</u> , eds. John Wang and Rodney Pearlman, New York:Plenum Press pps. 135-158, chapter 4, (1993)
*72	Reilly et al., "Oral delivery of antibodies: future pharmacokinetic trends" <u>Clin. Pharmacokinet.</u> 32(4):313-323 (1997)
*73	Sapan, "Immunoglobulin stability" <u>Biotechnol. Appl. Biochem.</u> 25:9-12 (1997)

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